

COMPOSITION AND METHOD FOR SUPPORTING CANCER TREATMENTS

Background of the Invention

1. Field of Invention

This invention relates generally to a novel composition to be used as a supporting composition in cancer treatments and more particularly to an herbal composition and method of using said herbal composition together with chemotherapy or radiation therapy (or both) in the treatment of cancer.

2. Description of Related Art

Normal cells grow and divide in an orderly and controlled manner. Cancer is a disease where cells become abnormal (cancerous cells) and begin to multiply without control to develop into an extra mass of tissue called a tumor. These cancerous cells can invade nearby tissues and spread through the blood stream and lymphatic system to other parts of the body.

Currently, the four primary types of cancer treatments are immunotherapy, surgery, radiation therapy, and chemotherapy. These cancer treatments may be applied alone or in conjunction with one another. Thus a cancer patient may undergo one or more treatments at a time. A single treatment would span a predefined period of time with therapies delivered at various timed intervals. Immunotherapy, also known as biological therapy or biological response modifier (BRM) therapy, tries to stimulate or restore the ability of the immune system to fight the disease. It is also used to lessen immune system related side effects that may be caused by some cancer treatments. Surgery seeks to directly remove the tumor from the body.

Radiation therapy, also known as radiotherapy, uses high-energy radiation from x-rays, gamma rays, neutrons, and other sources to kill cancer cells and shrink tumors by damaging the cells' genetic material. While cancerous cells are damaged permanently

and eventually die, some normal cells that are damaged in radiation therapy are also unable to repair themselves. Side effects that can occur during radiation therapy include skin irritation and hair loss in the area being treated and damage to the bone marrow.

Chemotherapy uses cytotoxic drugs, alone or in combination, to destroy cancer cells. Just as in radiation therapy, cancer cells can be damaged and eventually die. But only some healthy cells affected in the process can repair themselves after the chemotherapy. Cytotoxic drugs work by interfering with the ability of a growing cell to divide and reproduce itself. Thus, in addition to cancerous cells, other normal fast-dividing growing cells can also be affected. There can be an effect on blood cells forming in the bone marrow causing bone marrow suppression. There can also be an effect on cells in the digestive tract, in the lining of the mouth and in the reproductive system causing diarrhea and mouth soreness, and an effect on hair follicles causing hair loss.

Bone marrow suppression is one of the many side effects of chemotherapy and radiation therapy. It results in reduced blood cell production, including red blood cells, white blood cells, and platelets. Consequently, the patient will experience tiredness, from anemia, become more susceptible to infections, from leukopenia, and bruise easily and bleed more when getting a cut, from thrombocytopenia.

Drugs are used to counter the bone marrow suppression side effect. Epogen (Epoietin alpha) has been used to counter the side effect of anemia in cancer chemotherapy, and WinRho SD has been used to counter the side effect of thrombocytopenia.

Many of the treatments developed to coordinate with chemotherapy and radiation therapy to counter the side effect of leukopenia act on specific types of white blood cells, i.e. granulocytes, monocytes, and lymphocytes. Neupogen is a recombinant human granulocyte colony-stimulating factor (G-CSF) that stimulates the growth of neutrophils. Leukine is a recombinant human granulocyte-macrophage colony-stimulating factor that stimulates the production of neutrophils and macrophages. In animal laboratories and clinical trials, various interleukins, secreted by T-lymphocytes, have been used to stimulate productions of various white blood cells in the course of or after chemotherapy. interleukin 1 (IL-1) is responsible for B-cell and T-cell proliferation, interleukin 2 (IL-2) is responsible for the proliferation, growth, and activation of B-cells and T-cells,

interleukin 15 (IL-15) appears to be required for natural killer cells (NK cells) and CD8+ T-cells.

Herbs have also been found to have the activity of countering leukopenia side effect. For example, injection of extracts from *Sophora flavescens* roots has been reported to have reduced leukopenia side effect of chemotherapy and radiation therapy. Also, injection of *Uncaria tomentosa* water extracts in rats experiencing leukopenia from chemotherapy led to an increase in white blood cells.

Brief Summary of the Invention

The present invention is directed to a novel composition and method of using the novel composition in cancer treatments, preferably to reduce the bone marrow suppression side effect of such treatments. The novel composition is made of geranium oil and extractions from the root of Sophora plants, preferably *Sophora tonkinesis*, also known as *Sophora subprostrata*, (referred to herein as *Sophora tonkinesis*). The above "geranium oil" and "extractions from the root of Sophora plants" preferably refer to the main ingredients directly extracted from the oil and root respectively, but also includes main ingredients that are chemically synthesized or otherwise provided. The herbal composition can take on many forms e.g., powders, oil capsules, tablets, pills, liquid, syrup or pastes. The herbal composition can be made into and ingested as a food additive, dietary supplement, health food, decoction soup, or any other edible form. The herbal composition can be administered via various routes, i.e. oral, intravenous, or intraperitoneal, in specific dosages to mammalian animals undergoing chemotherapy or radiation therapy. For administration, the composition can be obtained by preparation, purchase, or any other means so one is in possession of the composition and administered before, during and after the cancer treatments.

Brief Description of the Drawings

Fig. 1 shows the compounds identified and their relative contents in the geranium oil produced in Kunming, China by the methods of gas chromatography/mass spectroscopy.

Fig. 2 shows the result of pharmacokinetics study of intravenous injection of matrine and matrine with and addition of geranium oil.

Fig. 3 shows the result of pharmacokinetics study of intravenous injection of oxymatrine and oxymatrine with the addition of geranium oil.

Detailed Description of the Invention

The present invention relates to a novel composition comprising geranium oil and extractions from the root of Sophora plants, preferably *Sophora tonkinesis*, and method of using the novel composition as a supporting drug or supplement in cancer treatments, preferably to reduce the bone marrow suppression side effect occurring in most of such treatments.

1. Geranium Oil

Geranium oil may be collected from steam distillation of the stem and leaves of the plant of division Magnoliophyta, class Magnoliopsida, order Geraniales, family Geraniaceae, and genus Pelargonium. Pelargoniums are native to South Africa and there are more than one hundred species in existence today, including hybridized garden species. Pelargoniums are now grown, and geranium oil is now produced, mainly in Algeria, Egypt, Morocco, Bourbon, China, and Australia. The present invention preferably uses geranium oil extracted from Pelargonium graveolens or Pelargonium roseum and Pelargonium terebinthineum grown in Kunming City of the Yunan Province in China. A gas chromatography/mass spectroscopy (GC-MS) result of the geranium oil produced in Kunming shows the constituent compounds and their relative contents (see Fig. 1). The generally known main constituents of geranium oil are citronellol, geraniol, geranyl formate, citronellyl formate, linalool, trans-rose oxide, and cis-rose oxide.

Certain specifications of geranium oil are set out in the National Standard of the People's Republic of China - GB 11959-89 which is incorporated herein by reference in their entirety, including any drawings. It adopts the same international standard of ISO 4731:1978 Oil of Geranium (Geranium Oil Standard). The Geranium Oil Standard specifies the outward characteristics of geranium oil, i.e. the geranium oil takes on a clear oil liquid form of a yellow greenish or amber color and has a distinct aroma. The same standard also specifies a relative density of 0.881 – 0.900 g/cm³, an optical rotation of -6°

to -14°, and a refractive index of 1.459 – 1.466 for geranium oil.

2. *Sophora tonkinesis*

The root of *Sophora tonkinesis* takes on a long curved tubular form with branches and is typically about 0.3 - 1.5 centimeters in diameter. The root is hardened and difficult to break. Its surface color ranges from grayish brown to suntan brown with longitudinal wrinkles and holes. The root has a bean scent and is extremely bitter. It is grown mainly in parts of China, i.e. the Guangdong province, Guangxi province, Guizhou province, Yunan province, and Jiangxi province.

The root contains 0.93% of alkaloids, of which 0.52% is matrine and 0.35% is oxymatrine. The other alkaloids identified in the root of *Sophora tonkinesis* are anagryne, methylcytisine, cytosine, sophocarpine, sophocarpine N-oxide, sophoramine, and sophoranol. The flavonic compounds identified in the root are sophoranone, sophoradin, sophoranochromene, sophoradochromene, pterocarpine, genistein, maackian, trifolirhizin, sitosterol, lu-peol, and a group of alkyl alcohol ester.

The principal alkaloid constituents of *Sophora tonkinesis* are also found in *Sophora alopecuroides*, *Sophora moorcroftiana*, and *Euchresta strigillosa*.

Result of pharmacokinetics study shows that in intravenous injections, the addition of geranium oil to matrine or oxymatrine will increase the absorption and metabolism of the respective compound (please see Fig. 2 and Fig. 3 for the changes in HPLC peak areas of matrine and matrine + geranium oil as time progresses). Furthermore, the composition of the present invention, containing oxymatrine, can also be taken orally to increase white blood cells. This is contrary to previously published data of animal experiments and clinical trials indicating that oxymatrine, when taken orally does not show any effect on increasing white blood cells, has to be injected through the muscles to increase white blood cells.

3. Composition

The composition can be formed into powders (composition powders) through the following steps. First, geranium oil and the root of *Sophora tonkinesis* are prepared separately. β -cyclodextrin is added to geranium oil to prevent evaporation, and excipients are added subsequently to form geranium oil powders. The geranium oil and

the excipients are about 31% and 69% by weight, respectively, of the geranium oil powders. Next, the root of *Sophora tonkinesis* is cut into thin pieces and then grounded. About 250 grams of the grounded *Sophora tonkinesis* root is mixed with 3000 ml of water, about 12 times the weight of the grounded root. The mixture is then boiled in a steam distillation bottle to heat and reflux for about 1 hour. Afterwards, the scum on the surface of the liquid is removed, and the liquid is filtered through a 100 mesh screen. The filtered liquid is then concentrated and about 66 grams of solid extracts of *Sophora tonkinesis* is obtained. Excipients are added to the solid extractions to form *Sophora tonkinesis* root powders. The *Sophora tonkinesis* extractions and the excipients are about 60% and 40% by weight, respectively, of the *Sophora tonkinesis* powders. Subsequently, the geranium oil powders and the *Sophora tonkinesis* root powders are mixed together with additional excipients to form the composition of the present invention into powder forms, wherein the geranium oil powders, *Sophora tonkinesis* root powders, and the excipients are about 55.94%, 0.958%, and 43.102% by weight, respectively, of the composition powders. The weight ratio of geranium oil and extractions of *Sophora tonkinesis* within the composition are about 30:1. The excipients to be used in the process to form powders can be starch, sugar spheres, fructose, sorbital crystalline etc. and those commonly used by one skilled in the art.

Alternatively, the geranium oil powders and the *Sophora tonkinesis* root powders can be mixed with glycerine and gelatin to form capsules. The composition can also be made into dietary supplement, health food (functional food), and food additives. One can also decoct the Pelargonium plant and Sophora roots to obtain a liquid form of the composition for direct oral intake as a medicine soup or for making into syrup or other forms of liquid composition. Sophora roots the Pelargonium plant can also be taken orally, in an edible form, separately at a timed interval.

EXAMPLE

Composition powders were administered orally to immunologically normal mice that were also given the 5-Fluorouracil (5-Fu) drug intraperitoneally.

The test substance, i.e. the composition powders, was prepared by dissolving the content in PBS.

Animals tested are 12 male BALB/c mice of 6-7 weeks old, weighing 22 ± 2 grams,

provided by Taiwan National University Medical Center Laboratory Animal Center. The animals are divided into two groups of 6 mice. Laboratory mice feeds manufactured by Purina (PMI5001) were used. Double-distilled water was provided for drinking. Laboratory mice wooden beddings manufactured by Beta Chip were used and changed 2-3 times weekly. Each group of 6 mice was kept in a feeding box of 29.2 x 19 x 12.7(cm). Micro-Isolator™ VCL Rack Housing System 70084A was used. Temperature and humidity were kept at $23\pm 2^{\circ}\text{C}$ and $60\pm 10\%$ respectively. The mice were given twelve hours of light and twelve hours of darkness.

Doses of 21mg and 7mg of test substance dissolved in PBS were administered to the two groups of test animals respectively, in a feeding volume of 0.2ml/mouse. The test substance were administered orally to the test animals the day after a single dose of the chemotherapeutic agent 5-Fu (135mg/kg, IP) was given and then once daily for the next nine (10 doses total) and thirteen consecutive days (14 doses in total) for the first and second group of mice respectively. On day 10, the first group of mice was sacrificed by anesthetizing with CO₂ and taking the blood from the heart to determine the cell counts of erythrocytes (RBC), platelets (PLT), total leukocytes (WBC), and differential leukocytes counts: lymphocytes (LY), monocytes (MO), and granulocytes (GR). On day 14, the second group of mice was sacrificed in the same manner to determine the same blood cell counts. The control employed in the experiment were normal mice without any injections.

As shown in the table below, 7mg/mouse of test substance had the significant effect of increasing the number of red blood cells(RBC) and preventing the reduction of the number of WBC, LY, MO, and GR in mice injected with 5-Fu. The effect was more pronounced with the dosage of 7mg/mouse. On day 10, the average WBC count of normal mice was $6.94 \pm 1.647 \times 10^3/\mu\text{l}$, and the mice treated with 5-Fu had an average WBC count of $4.17 \pm 0.677 \times 10^3/\mu\text{l}$. On the other hand, mice treated with 7mg/mouse of test substance and 5-Fu had an average WBC count of $6.24 \pm 1.924 \times 10^3/\mu\text{l}$, showing only 25% of the bone marrow suppression effect of 5-Fu. Differential leukocyte count showed that the suppression effect with respect to lymphocytes in test animals treated with 7mg/mous of test substance and 5-Fu was only 12% of that of the test animals treated with 5-Fu only. With respect to monocytes, the suppression effect in test animals treated with 7mg/mouse of test

substance and 5-Fu was only 21 % of that of test animals treated with 5-Fu. With respect to granulocytes, the suppression effect in test animals treated with 7mg/mouse of test substance and 5-Fu was 46% of that of test animals treated with 5-Fu. On day 14, the total leukocyte and differential leukocyte counts of mice treated with 7mg/mouse of test substance and 5-Fu continued to increase to a higher level than that of mice treated with 5-Fu only.

Effect of 「geranium oil + *Sophora tonkinesis* extractions」 on the side effects of reduction in blood cell counts caused by 5-Fu -

	Normal	5-Fu	21 mg/mouse S. tonkinesis /5-Fu	7 mg/mouse S. tonkinesis /5-Fu
Day 10				
RBC ($10^6/\mu\text{l}$)	9.09 ± 0.137	7.86 ± 0.171	7.66 ± 0.316	$8.52 \pm 0.627^*$
PLT ($10^3/\mu\text{l}$)	990 ± 65.7	2828 ± 632.4	2441 ± 441.4	2099 ± 731.5
WBC ($10^3/\mu\text{l}$)	6.94 ± 1.647	4.17 ± 0.677	4.63 ± 0.772	$6.24 \pm 1.924^*$
LY ($10^3/\mu\text{l}$)	5.30 ± 1.369	3.66 ± 0.648	4.15 ± 0.538	$5.10 \pm 1.261^*$
MO ($10^3/\mu\text{l}$)	0.39 ± 0.035	0.25 ± 0.046	0.26 ± 0.154	$0.36 \pm 0.131^*$
GR ($10^3/\mu\text{l}$)	1.24 ± 0.284	0.25 ± 0.050	0.22 ± 0.104	$0.78 \pm 0.559^*$
Day 14				
RBC ($10^6/\mu\text{l}$)	9.76 ± 0.269	8.09 ± 0.331	8.19 ± 0.160	8.23 ± 0.326
PLT ($10^3/\mu\text{l}$)	985 ± 216.5	2219 ± 750.2	2461 ± 195.4	2309 ± 687.5
WBC ($10^3/\mu\text{l}$)	8.03 ± 1.408	7.98 ± 1.575	7.70 ± 0.599	8.48 ± 2.052
LY ($10^3/\mu\text{l}$)	6.53 ± 1.470	5.75 ± 0.880	6.10 ± 0.397	6.56 ± 1.591
MO ($10^3/\mu\text{l}$)	0.35 ± 0.092	0.59 ± 0.316	0.39 ± 0.124	0.44 ± 0.140
GR ($10^3/\mu\text{l}$)	1.15 ± 0.243	1.65 ± 0.756	1.21 ± 0.353	1.47 ± 0.560

1. Results are expressed in mean \pm standard deviation (mean \pm SD).

2. The experimental group and the 5-Fu group are compared using Dunnett's t-test, "*" means $p < 0.05$, "**" means $p < 0.01$, and "***" means $p < 0.001$.

The weight of mice treated with 7mg/mouse and 21mg/mouse decreased slightly, as the days progresses, as compared to the normal mice.

Effect of 「geranium oil + *Sophora tonkinesis* extractions」 on the side effects of weight change caused by 5-Fu

	Day -2	Day 0	Day 6	Day 10	Day 14
Normal control	20.3 ± 1.90 (n=18)	21.6 ± 1.85 (n=18)	23.1 ± 1.94^a (n=18)	24.0 ± 1.89^a (n=12)	24.6 ± 2.21 (n=6)
5-Fu	20.7 ± 0.90 (n=12)	22.0 ± 0.93 (n=12)	21.9 ± 1.20^{ab} (n=12)	22.3 ± 1.53^b (n=12)	23.5 ± 0.78 (n=6)
G-CSF/5-Fu	20.5 ± 1.76 (n=12)	22.0 ± 1.84 (n=12)	22.0 ± 1.84^{ab} (n=12)	22.5 ± 1.78^{ab} (n=12)	24.0 ± 1.64 (n=6)

21 mg/mouse	19.6 ± 1.42	21.2 ± 1.38	21.4 ± 1.71^b	21.4 ± 1.80^b	22.3 ± 1.57
S.	(n=12)	(n=12)	(n=12)	(n=12)	(n=6)
<i>tonkinesis</i> /5-Fu					
7 mg/mouse	19.8 ± 1.50	21.4 ± 1.97	22.2 ± 1.67^{ab}	22.6 ± 1.68^{ab}	23.1 ± 3.04
S.	(n=12)	(n=12)	(n=12)	(n=12)	(n=6)
<i>tonkinesis</i> /5-Fu					

1. Results are expressed in mean \pm standard deviation (mean \pm SD).
2. At the same moment in time, Duncan's statistical analysis is used among the groups. Different alphabets stands for significant differences ($p < 0.05$).

In comparison with another type of treatment relating to the reduction of bone marrow suppression using G-CSF, the bone marrow suppression effect was not as significantly reduced as that of the tested novel composition of the present invention. On day 10, the normal mice's average WBC count was $6.94 \pm 1.647 \times 10^3/\mu\text{l}$, and the mice treated with 5-Fu had an average WBC count of $4.17 \pm 0.677 \times 10^3/\mu\text{l}$. On the other hand, mice treated with $135\mu\text{g}/\text{mouse}$ of G-CSF and 5-Fu had an average WBC count of $5.46 \pm 2.338 \times 10^3/\mu\text{l}$, showing 51% of the bone marrow suppression effect of 5-Fu. Differential leukocyte count showed that the suppression effect with respect to lymphocytes in test animals treated with $135\mu\text{g}/\text{mouse}$ of G-CSF and 5-Fu was only 18% of that of the test animals treated with 5-Fu only. With respect to monocytes, the suppression effect in test animals treated with $135\mu\text{g}/\text{mouse}$ of G-CSF and 5-Fu was only 7 % of that of test animals treated with 5-Fu. With respect to granulocytes, the suppression effect in test animals treated with $135\mu\text{g}/\text{mouse}$ of G-CSF and 5-Fu was 54% of that of test animals treated with 5-Fu. On day 14, only the total leukocyte count and differential leukocyte count, with respect to lymphocytes of mice treated with $135\mu\text{g}/\text{mouse}$ of G-CSF and 5-Fu, continued to increase to a higher level than that of mice treated with 5-Fu only.

Effect of G-CSF on the reduction of 5-Fu's side effect- change in blood cell counts

	Normal	5-Fu	$135\mu\text{g}/\text{mouse}$ C-GSF/5-Fu
Day 10			
RBC ($10^6/\mu\text{l}$)	9.09 ± 0.137	7.86 ± 0.171	7.82 ± 0.424
PLT ($10^3/\mu\text{l}$)	990 ± 65.7	2828 ± 632.4	2303 ± 491.7
WBC ($10^3/\mu\text{l}$)	6.94 ± 1.647	4.17 ± 0.677	5.46 ± 2.338
LY ($10^3/\mu\text{l}$)	5.30 ± 1.369	3.66 ± 0.648	$5.01 \pm 1.372^*$
MO ($10^3/\mu\text{l}$)	0.39 ± 0.035	0.25 ± 0.046	$0.38 \pm 0.141^*$

GR ($10^3/\mu\text{l}$)	1.24 ± 0.284	0.25 ± 0.050	$0.71 \pm 0.268^{**}$
Day 14			
RBC ($10^6/\mu\text{l}$)	9.76 ± 0.269	8.09 ± 0.331	8.02 ± 0.340
PLT ($10^3/\mu\text{l}$)	985 ± 216.5	2219 ± 750.2	2105 ± 378.1
WBC ($10^3/\mu\text{l}$)	8.03 ± 1.408	7.98 ± 1.575	8.34 ± 1.454
LY ($10^3/\mu\text{l}$)	6.53 ± 1.470	5.75 ± 0.880	6.12 ± 1.164
MO ($10^3/\mu\text{l}$)	0.35 ± 0.092	0.59 ± 0.316	0.58 ± 0.266
GR ($10^3/\mu\text{l}$)	1.15 ± 0.243	1.65 ± 0.756	1.64 ± 0.405

1. Results are expressed in mean \pm standard deviation (mean \pm SD).

2. The experimental group and the 5-Fu group are compared using Dunnett's t-test, "*" means $p < 0.05$, "***" means $p < 0.01$, and "****" means $p < 0.001$.

The weight of the mice shows no significant difference among the groups.

Effect of G-CSF on the reduction of 5-Fu's side effect- weight change

	Day -2	Day 0	Day 6	Day 10	Day 14
Normal control	20.3 ± 1.90 (n=18)	21.6 ± 1.85 (n=18)	23.1 ± 1.94 (n=18)	24.0 ± 1.89^a (n=12)	24.6 ± 2.21 (n=6)
5-Fu	20.7 ± 0.90 (n=12)	22.0 ± 0.93 (n=12)	21.9 ± 1.20 (n=12)	22.3 ± 1.53^b (n=12)	23.5 ± 0.78 (n=6)
G-CSF/5-Fu	20.5 ± 1.76 (n=12)	22.0 ± 1.84 (n=12)	22.0 ± 1.84 (n=12)	22.5 ± 1.78^b (n=12)	24.0 ± 1.64 (n=6)

1. Results are expressed in mean \pm standard deviation (mean \pm SD).

2. At the same moment in time, Duncan's statistical analysis is used among the groups.

Different alphabets stands for significant differences ($p < 0.05$).

The composition of *Sophora tonkinesis* and geranium oil does in fact significantly reduces the bone marrow suppression effect of 5-Fu and is performing better even than the G-CSF treatment. The ability of the composition of the present invention to reduce bone marrow suppression effect makes it a good candidate as a supporting drug or supplement to be used in cancer treatments that induce bone marrow suppression. In particular, the composition of the present invention may be used with chemotherapy and or radiation therapy to increase the leukocyte count. For example, the composition of the present invention may be used with 5-Fu, doxorubicin and other chemotherapeutic agents just as Neupogen is also used with 5-Fu as well as doxorubicin and many other types of chemotherapy to stimulate the growth of neutrophils whose number is originally reduced by chemotherapy.

Human dosages can be calculated based on the dosages used for mice in the experiment. In accordance with accepted clinical trial practice, mice dosages are divided by a factor of 10 in order to obtain suitable and safe dosages for human. A range of dosage is calculated based on the premise that the mice weighed about 20-25 grams. The calculation is shown as follows.

$$7\text{mg/mouse/day} \rightarrow 7\text{mg}/20\text{g/day} \times 50 \rightarrow 350\text{mg/kg/day} \div 10 \times 60 = 2100\text{mg}/60\text{kg/day}$$

$$7\text{mg/mouse/day} \rightarrow 7\text{mg}/25\text{g/day} \times 50 \rightarrow 280\text{mg/kg/day} \div 10 \times 60 = 1680\text{mg}/60\text{kg/day}$$

$$21\text{mg/mouse/day} \rightarrow 21\text{mg}/20\text{g/day} \times 50 \rightarrow 1050\text{mg/kg/day} \div 10 \times 60 = 6300\text{mg}/60\text{kg/day}$$

$$21\text{mg/mouse/day} \rightarrow 21\text{mg}/25\text{g/day} \times 50 \rightarrow 840\text{mg/kg/day} \div 10 \times 60 = 5040\text{mg}/60\text{kg/day}$$